Interagency Report Control No 0180-DOA-AN

United States Department of Agriculture Animal and Plant Health Inspection Service CONTINUATION SHEET FOR ANNUAL REPORT OF RESEARCH FACILITY

1. Registration No. 84-F-0001

FORM APPROVED NO. 0579-0036

OMB

2. HEADQUARTERS RESEARCH FACILITY (Name and Address, as registered with

USDA including zip code) USDA, APHIS, WS, NWRC 4101 LaPorte Avenue Fort Collins, CO 80521 (970) 266-6000

	B. Number of	C. Number of	D. Number of animals	heets if necessary or use APHIS FORM 7023A) E. Number of animals upon which teaching	T
A. Animals Covered By The Animal Welfare Regulations 12. &/OR 13. Other (List by Species)	animals being bred, conditioned, or held for use in teaching, testing, experiments, research, or surgery but not yet used for such purposes.	animals upon which teaching, research, experiments, or tests were conducted involving no pain, distress, or use of pain- relieving drugs	upon which experiments, teaching, research, surgery, or tests were conducted involving accompanying pain or distress to the animals and for which appropriate anesthetic, analgesic, or tranquilizing drugs were used.	experiments, research, surgery, or tests were conducted involving accompanying pain or distress to the animals and for which the use of appropriate anesthetic, analgesic, or tranquilizing drugs would have adversely affected the procedures, results, or interpretation of the teaching, research, experiments, surgery, or tests. (An explanation of the procedures producing pain or distress in those animals and the reasons such drugs were not used must be attached to this report.	Total No. of Animals (Cols. C + D + E)
Coyotes	39	102	24	8	134
Raccoons	0	54	0	0	54
Polynesian Rats	142	141	0	0	141
Roof Rats	130	124	0	0	124
Norway Rats	55	35	0	0	35
Gambian Pouched Rat	32	0	0	0	0
House Mouse	163	175	0	0	175
Voles	39	0	0	0	0
Grey Squirrel	0	44	0	0	44
Pocket Gopher	21	0	0	0	0
Prairie Dogs	0	91	0	18	109
Big Brown Bats	36	0	0	0	0
Beaver	33	13	0	0	13
and following actual research, teaching, to principal investigator has considered alter facility is adhering to the standards and reinvestigator and approved by the Institution the IACUC-approved exceptions, this sure 4) The attending veterinarian for this researment care and use. CERTIF	esting, surgery, or a mattives to painful egulations under the onal Animal Care an mmary includes a bearch facility has ap	experimentation we procedures. e Act, and it has re nd Use Committee rief explanation of to propriate authority Y HEADQU	are followed by this research quired that exceptions to th (IACUC). A summary of al the exceptions, as well as th to ensure the provision of a	oriate use of anesthetic, analgesic, and tranquilizing drug of facility. e standards and regulations be specified and explained is such exceptions is attached to this annual report in additional expectations and the species and number of animals affected, dequate veterinary care and to oversee the adequacy of ARCH FACILITY OFFICIAL	2) Each 3) This by the principal lition to identifying
(Chief Executive Officer or Leg			correct, and complete	e (7 U.S.C. Section 2143). of CEO or Institutional Official (type or print)	Date Signed
(b)(6),(b)(7)(c)				(b)(6),(b)(7)(c)	11/30/

Column E Explanation - QA 1440

1. Registration Number: 84-F-0001

2. Number of animals used in this study during this reporting period: Nine (9)

3. Species (common name) of animals used in study: Canis latrans (coyote)

- 4. Explain procedure producing pain and/or distress: Coyotes consumed a bait that contained theobromine and caffeine
- 5. Provide scientific justification why pain or distress could not be relieved. State method or means used to determine that pain and/or distress relief would interfere with test results. (For Federally mandated testing, see item 6 below)

This was an efficacy test for a toxicant. Co-administration of drugs may impact the observed effects of the test materials and nullify the experimental conclusions. Additionally, only minor distress was observed during this study.

6. What, if any federal regulations require this procedure?

Agency: US Environmental Protection Agency

CFR: Title 40, FIFRA

Column E Explanation QA-1446

- 1. Registration Number: 84-F-0001
- 2. Number of animals used in this study during this reporting period: 18
- 3. Species (common name) of animals used in study: Black-tailed prairie dog
- 4. Explain procedure producing pain and/or distress: Animals were gavaged with a predetermined amount of chlorophacinone (an anticoagulant) technical material in propylene glycol. The gavage procedure itself did not produce more than temporary stress for the animal. However, the effects of the anticoagulant itself led to more than temporary distress in some animals.
- 5. Provide scientific justification why pain or distress could not be relieved. State method or means used to determine that pain and/or distress relief would interfere with test results. (For Federally mandated testing, see item 6 below): Of the 70 animals used, 18 animals were considered to be experiencing more than momentary pain or distress. Because the Environmental Protection Agency requires death as an endpoint for toxicants, only those animals suffering extreme pain or distress that were unlikely to survive were euthanized. Of the 20 animals that died during the course of the study, four animals were euthanized. One animal that appeared highly likely to die recovered at the end of the study, illustrating the difficulty in balancing animal welfare issues with obtaining scientifically valid data.
- 6. What, if any federal regulations require this procedure?

Agency: Environmental Protection Agency

CFR: U.S. EPA 1996. Ecological Effects Test Guidelines: Wild Mammal Acute Toxicity. OPPTS 850.2400 and OPP Pesticide Assessment Subdivision G: Product Performance. Section 96-12: Rodenticides on Farm and Rangeland.